

## CLAIMS

1. A fusion protein having epitopes of at least two of the autoantigens glutamic acid decarboxylase (GAD65), islet cell antigen (IA2) and preproinsulin (PPINS) wherein said epitopes are connected with a linker peptide, said fusion  
 5 protein being able to bind to a solid phase.

2. The fusion protein according to claim 1 having epitopes of each of the autoantigens GAD65, IA2 and PPINS.

3. The fusion protein according to claim 2 wherein  
 - the epitope of IA2 comprises the amino acids 771-979 of  
 10 the amino acid sequence shown in Figure 2a,  
 - the epitope of GAD65 comprises the amino acids 102-585 of the amino acid sequence shown in Figure 2b, and  
 - the epitope of PPINS comprises all the amino acids 1-110 of the amino acid sequence shown in Figure 2c.

15 4. The fusion protein according to claim 1 wherein the linker peptide comprises lysine and argine residues.

5. The fusion protein according to claim 4 wherein said linker peptide is provided with a member of an affinity binding pair so as to enable the binding of said fusion  
 20 protein to the solid phase.

6. The fusion protein according to claim 5 wherein the affinity binding pair is biotin - streptavidin.

7. A cDNA encoding the fusion protein according to claim 1 wherein said cDNA comprises the nucleotide sequences  
 25 encoding the epitopes of at least two of the autoantigens glutamic acid decarboxylase (GAD65), islet cell antigen (IA2) and preproinsulin (PPINS).

8. A cDNA encoding the fusion protein according to claim 3

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wherein said cDNA comprises the nucleotide sequences  
 a) nucleotides 1311 to 1755 of the sequence according to  
 Figures 3a to 3b encoding GAD65, aa 102-585,  
 b) nucleotides 2313 to 2937 of the sequence according to  
 Figures 3c to 3e encoding IA2, aa 771-979, and  
 c) nucleotides 2424 to 2610 and 3397 to 3539 of the  
 sequence according to Figure 3f-3i encoding PPINS, aa 1-  
 110, where said nucleotide sequences a), b) and c) can  
 appear in any relative order.

10 9. A vector comprising the cDNA according to claim 7 ~~or 8.~~

10. An E. coli cell encompassing the cDNA according to  
 claim 7.

11. An immunoassay for the simultaneous determination in a  
 sample of a person's body fluid of at least two insulin  
 15 dependent diabetes mellitus (IDDM) related autoantibodies,  
 wherein each autoantibody is specific for an epitope of the  
 autoantigens glutamic acid decarboxylase (GAD65), islet  
 cell antigen (IA2) or preproinsulin (PPINS), said  
 immunoassay comprising the steps of  
 20 - incubating said sample with a fusion protein according to  
 claim 1, said fusion protein being bound to a solid  
 support,  
 - adding at least one labeled reagent capable of binding to  
 one or more of said autoantibodies, and  
 25 - quantifying the signals from the labels bound to the  
 solid phase.

12. The immunoassay according to claim 11 wherein the  
 labeled reagent is an anti-human monoclonal antibody.

13. The immunoassay according to claim 11 wherein the  
 30 labeled reagent comprises at least two antigens labeled  
 with different labels, each antigen being one of the  
 autoantigens GAD65, IA2 or PPINS; or proteins comprising  
 epitopes thereof.

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14. The immunoassay according to claim 11 wherein the  
 labeled reagent comprises three antigens labeled with the  
 same label, each antigen being one of the autoantigens  
 5 GAD65, IA2 or PPINS; or proteins comprising epitopes  
 thereof.

15. The immunoassay according to claim 11 wherein the label  
 is a fluorescent lanthanide chelate.

16. A method for diagnosing a person's risk of developing  
 10 insulin dependent diabetes mellitus (IDDM), said method  
 comprising the determination in a sample of said person's  
 body fluid of at least two insulin dependent diabetes  
 mellitus (IDDM) related autoantibodies specific for an  
 epitope of the autoantigens glutamic acid decarboxylase  
 15 (GAD65), islet cell antigen (IA2) or preproinsulin (PPINS),  
 wherein the presence of at least two of said autoantibodies  
 are indicative for said person's risk of developing IDDM.

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